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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/760,384	01/11/2001	Hau H. Duong	A-68718-2/RFT/RMS/RMK	2482
32940	7590 08/24/2004		EXAMINER	
DORSEY & WHITNEY LLP INTELLECTUAL PROPERTY DEPARTMENT			SINES, BRIAN J	
4 EMBARCA	ADERO CENTER	ACTIVILIA I	ART UNIT	PAPER NUMBER
SUITE 3400	SCO, CA 94111		1743	
BANTKAIN	015CO, CA 94111		DATE MAILED: 08/24/2004	

Please find below and/or attached an Office communication concerning this application or proceeding.

	Application No.	Applicant(s)	Mu-
	09/760,384	DUONG ET AL.	
Office Action Summary	Examiner	Art Unit	
	Brian J. Sines	1743	
The MAILING DATE of this communication Period for Reply	appears on the cover sheet w	ith the correspondence address	
A SHORTENED STATUTORY PERIOD FOR RETHE MAILING DATE OF THIS COMMUNICATION - Extensions of time may be available under the provisions of 37 CF after SIX (6) MONTHS from the mailing date of this communication. If the period for reply specified above is less than thirty (30) days, If NO period for reply is specified above, the maximum statutory period for reply within the set or extended period for reply will, by some Any reply received by the Office later than three months after the rearned patent term adjustment. See 37 CFR 1.704(b).	DN. FR 1.136(a). In no event, however, may a in. a reply within the statutory minimum of thire eriod will apply and will expire SIX (6) MON tatute, cause the application to become Af	reply be timely filed by (30) days will be considered timely. ITHS from the mailing date of this communication ANDONED (35 U.S.C. § 133)	on.
Status			
1) Responsive to communication(s) filed on 2	7/2/2004.		
	This action is non-final.		
3) Since this application is in condition for all	owance except for formal matt	ers, prosecution as to the merits i	s
closed in accordance with the practice und	ler <i>Ex par</i> te Quayle, 1935 C.D	. 11, 453 O.G. 213.	
Disposition of Claims			
4) Claim(s) 24-42 is/are pending in the applic	ation.		
4a) Of the above claim(s) is/are with	drawn from consideration.		
5) Claim(s) is/are allowed.			
6)⊠ Claim(s) <u>24-42</u> is/are rejected.			
7) Claim(s) is/are objected to.	•		
8) Claim(s) are subject to restriction ar	nd/or election requirement.		
Application Papers			
9)☐ The specification is objected to by the Exam	niner.		
10)☐ The drawing(s) filed on is/are: a)☐	accepted or b) objected to	by the Examiner.	
Applicant may not request that any objection to	the drawing(s) be held in abeyar	ce. See 37 CFR 1.85(a).	
Replacement drawing sheet(s) including the co		-	d).
11) The oath or declaration is objected to by the	e Examiner. Note the attached	Office Action or form PTO-152.	
Priority under 35 U.S.C. § 119			
12) Acknowledgment is made of a claim for fore a) All b) Some * c) None of:		119(a)-(d) or (f).	
1. ☐ Certified copies of the priority docum			
2. Certified copies of the priority docum			
3. Copies of the certified copies of the		received in this National Stage	
application from the International Bu * See the attached detailed Office action for a		received	
oso the attached detailed office action for a	ist of the certified copies flot	received.	
attachment(s)			
) Notice of References Cited (PTO-892)	4) Interview S	ummary (PTO-413)	
) Notice of Draftsperson's Patent Drawing Review (PTO-948)) Information Disclosure Statement(s) (PTO-1449 or PTO/SB	Paper Nots)/Mail Date	
	_	formal Patent Application (PTO-152)	

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DETAILED ACTION

Claim Rejections - 35 USC § 112

The following is a quotation of the second paragraph of 35 U.S.C. 112:

The specification shall conclude with one or more claims particularly pointing out and distinctly claiming the subject matter which the applicant regards as his invention.

Claims 30 – 40 are rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention.

Claim 30 recites the limitation "said assay complexes" in line 2. There is insufficient antecedent basis for this limitation in the claim.

Claim Rejections - 35 USC § 102

The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless -

(b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.

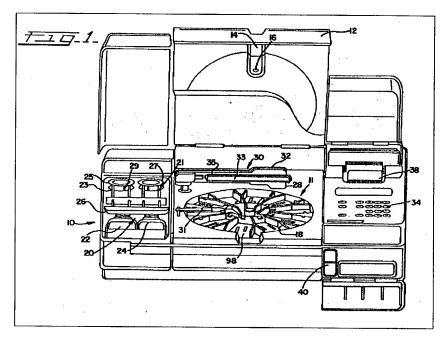
Claims 24 – 26, 41 and 42 are rejected under 35 U.S.C. 102(b) as being anticipated by Holen et al. (U.S. Pat. No. 5,320,808 A). Holen et al. teach a semi-automated sample analyzer and subsystems for simultaneously performing a plurality of immunoassays utilizing reaction cartridges. A carousel is provided to position and hold a plurality of reaction cartridges. Each cartridge includes a plurality of isolated test sites formed in a two-dimensional array in a solid-phase binding layer contained within a reaction well, which is adapted to contain a biological sample to be assayed (see Abstract).

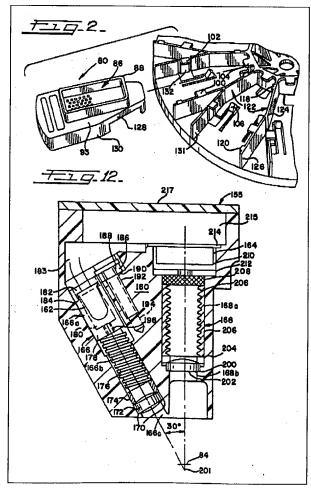
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Regarding claims 24, 41 and 42, Holen et al. anticipate the recited method of analyzing a plurality of biochips, wherein the method is comprising the steps of:

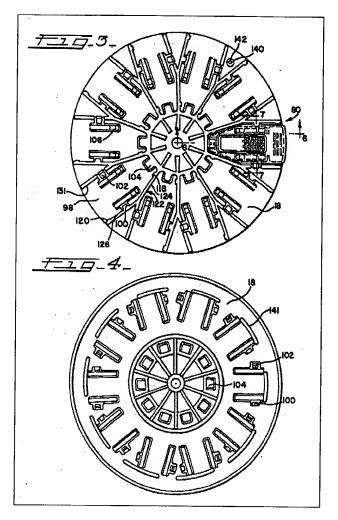
- (a) inserting a first biochip (reaction cartridge 80) into a first station (opening 98 on carousel 18) of an analysis device (10);
- (b) inserting a second biochip into a second station of the analysis device, wherein each of the first and second biochips comprise a substrate (test card 82) comprising an array comprising a plurality of test sites (84), wherein each test site (84) is comprising:
- i) a different capture binding ligand (e.g., a capture reagent, such as antibodies, antigens, anti-biotin, avidin, lectins, peptide sequence probes or specific allergen which binds human IgE class antibodies from a patient serum sample);
- ii) a different target analyte (e.g., a specific binding component, such as antigens from a patients sample, such as from serum, blood, urine, cerebrospinal fluid (CSF) or saliva); and
- iii) a label (e.g., a human IgE class antibody conjugated to an enzyme, such as alkaline phosphatase or horse radish peroxide (HRPO) conjugate, or any other detectable enzymatic or fluorogenic conjugate);
- (c) detecting the presence of the label on the first biochip utilizing optical reader (32) from the test sites (84) from the reaction cartridge (80); and
- (d) detecting the presence of the label on the second biochip (see col. 6, line 10 col. col. 20, line 38; figures 2, 3 & 5).

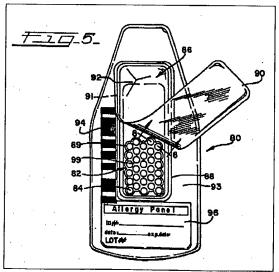
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Regarding claims 25 and 26, Holen et al. anticipate that the rotatably-mounted optical reader arm (35) is moved over each reaction cartridge (80). Holen et al. teach that under microprocessor control, the carousel (18) and optical reader arm (35) move in cooperation to sequentially position the optical reader (32) over each selected test site (84) until all selected test sites have been read (see col. 8, lines 4-35; col. 17, line 26-col. 18, line 66).

Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.

The factual inquiries set forth in *Graham* v. *John Deere Co.*, 383 U.S. 1, 148 USPQ 459 (1966), that are applied for establishing a background for determining obviousness under 35 U.S.C. 103(a) are summarized as follows:

- 1. Determining the scope and contents of the prior art.
- 2. Ascertaining the differences between the prior art and the claims at issue.
- 3. Resolving the level of ordinary skill in the pertinent art.
- 4. Considering objective evidence present in the application indicating obviousness or nonobviousness.
- 1. Claims 27 29 are rejected under 35 U.S.C. 103(a) as being unpatentable over Holen et al. Regarding claim 27, Holen et al. do not specifically teach the incorporation of a plurality of detectors within the system. Holen et al. do teach the incorporation of a single optical reader or detector in determining assay results (see e.g., col. 8, lines 4 35). The Courts have held that the mere duplication of parts, without any new or unexpected results, is within the ambit of one of ordinary skill in the art. See *In re Harza*, 124 USPQ 378 (CCPA 1960) (see MPEP § 2144.04).

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Therefore, it would have been obvious to a person of ordinary skill in the art to incorporate a plurality of detectors within the analyzer system, as taught by Holen et al., to facilitate the detection and analysis of the plurality of reaction cartridges used within the Holen et al. system. Regarding claim 28, Holen et al. do teach the use of fluorescence detection (see col. 19, line 65 – col. 20, line 17; col. 36, line 28 – col. 37, line 19). Regarding claim 29, Holen et al. teach the use of an electronic detector, e.g., a solid-state device, i.e., light-emitting diode (LED) (see col. 20, lines 25 – 38).

2. Claims 30, 31, 33 and 37 are rejected under 35 U.S.C. 103(a) as being unpatentable over Holen et al. in view of Kearney et al. (U.S. Pat. No. 5,759,777 A). Regarding claims 30 and 31, Holen et al. do not specifically teach that the capture binding ligands are nucleic acid probes, the target analytes are target nucleic acid sequences and the assay complexes are hybridization complexes. However, Holen et al. do indicate that their system can be utilized for DNA hybridization capacity studies (see col. 13, lines 34 - 57). Kearney et al. do teach the use of nucleic acid hybridization assays for the sensitive and specific detection, and isolation of specific nucleic acid sequences in clinical samples (see col. 1, lines 1-61). Consequently, a person of ordinary skill in the art would accordingly have had a reasonable expectation of success of incorporating the teachings of Kearney et al. with the system of Holen et al. The Courts have held that the prior art can be modified or combined to reject claims as prima facie obvious as long as there is a reasonable expectation of success. See In re Merck & Co., Inc., 800 F.2d 1091, 231 USPQ 375 (Fed. Cir. 1986) (see MPEP § 2143.02). Therefore, it would have been obvious to a person of ordinary skill in the art to incorporate the teachings of Kearney et al. with the system of Holen et al., in order to effectively perform hybridization assays. Regarding claim 33,

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Kearney et al. do teach the use of labeled probes, which inherently indicate hybridization during detection (see col. 7, lines 56 - 59). Regarding claim 37, Kearney et al. teach the use of fluorescent labels, such as fluoroscein (see col. 10, lines 1 - 15).

3. Claim 32 is rejected under 35 U.S.C. 103(a) as being unpatentable over Holen et al. in view of Kearney et al., as applied to claims 30, 31, 33 and 37 above, and further in view of Yabusaki et al. (U.S. Pat. No. 4,599,303). Holen et al. and Kearney et al. do not specifically teach the use of labels covalently attached to the target sequences. Yabusaki et al. do teach a method wherein probes or labels are covalently attached to the target nucleic acid sequence. Yabusaki et al. do teach a method of identifying specific nucleic acid sequences in which the target nucleic acid sequence is reacted with a probe or label under conditions where hybridization of the probe with the target sequence will occur. Following hybridization, the sample is subjected to a photochemical or chemical procedure, which causes covalent crosslinking of the probe to the target complementary sequence. Following cross-linking, the uncrosslinked probe is separated from the covalently crosslinked probe-target complex (see col. 2, lines 24 - 68; col. 3, lines 1 - 23). As evidenced by Yabusaki et al., an artisan of ordinary skill in the art would accordingly have recognized the suitability of incorporating this disclosed methodology of detecting target nucleic acid sequences with the methodology of Holen et al. in view of Kearney et al. for the intended purpose of providing for an effective means for facilitating the sensitive detection of extremely low concentrations of specific nucleic acid base sequences (see MPEP § 2144.07). Furthermore, as disclosed by Yabusaki et al., an artisan of ordinary skill in the art would accordingly have had a reasonable expectation of success of incorporating such a methodology for facilitating the effective detection of target nucleic acid

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sequences. The Courts have held that the prior art can be modified or combined to reject claims as *prima facie* obvious as long as there is a reasonable expectation of success. See *In re Merck & Co., Inc.*, 800 F.2d 1091, 231 USPQ 375 (Fed. Cir. 1986) (see MPEP § 2143.02). therefore, it would have been obvious to a person of ordinary skill in the art to incorporate the method disclosed by Yabusaki et al., with the method of Holen et al. and Kearney et al. in order to provide an effective means for detecting target nucleic acid sequences.

- 4. Claim 34 is rejected under 35 U.S.C. 103(a) as being unpatentable over Holen et al. in view of Kearney et al., as applied to claims 30, 31, 33 and 37 above, and further in view of Batz et al. (U.S. Pat. No. 6,117,973 A). Holen et al. and Kearney et al. do not specifically teach the use of intercalators. Batz et al. do teach the use of intercalators as probes or labels in hybridization assays (see col. 3, line 12 col. 4, line 8; col. 8, lines 4 46). The Courts have held that the selection of a known material, based upon its suitability for the intended use, is within the ambit of one of ordinary skill in the art. See *In re Leshin*, 125 USPQ 416 (CCPA 1960) (see MPEP § 2144.07). Therefore, it would have been obvious to an artisan of ordinary skill in the art to incorporate the use of an intercalator, as taught by Batz et al., with the methodology of Holen et al. and Kearney et al. in order to provide for an effective means of indicating hybridization.
- 5. Claims 35, 36, 38 and 39 are rejected under 35 U.S.C. 103(a) as being unpatentable over Holen et al. in view of Kearney et al., as applied to claims 30, 31, 33 and 37 above, and further in view of Meade et al. (U.S. Pat. No. 5,780,234 A). Regarding claims 35, 36 and 38, Meade et al. teach hybridization complexes essentially comprising: capture probes hybridized to a first domain of a target sequence; and label probes, e.g., covalently attached electron transfer

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moieties, hybridized to a second domain of the target sequence (see col. 5, line 29 - col. 6, line 62). Regarding claims 38 and 39, Holen et al. do not specifically teach the use of electron transfer moieties, such as transition metal complexes, as labels. Meade et al. do teach the use of electron transfer moieties, such as transtion metal complexes as labels or probes for assays (see col. 7, lines 14-67; col. 8, lines 1-67; col. 10, lines 44-59). Hence, as evidenced by Meade et al., a person of ordinary skill in the art would have recognized the suitability of using the teachings of Meade et al. regarding the use of electron transfer moieties and transition metal complexes as labels or probes for the intention of performing hybridization assays (see MPEP § 2144.07). The Courts have held that the selection of a known material, based upon its suitability for the intended use, is within the ambit of one of ordinary skill in the art. See In re Leshin, 125 USPQ 416 (CCPA 1960) (see MPEP § 2144.07). Furthermore, Meade et al. do teach that their disclosed technique is suited for automated probe assays (see col. 8, lines 62-67). Consequently, a person of ordinary skill in the art would accordingly have had a reasonable expectation of success of incorporating the teachings of Meade et al. with the system of Holen et al. The Courts have held that the prior art can be modified or combined to reject claims as prima facie obvious as long as there is a reasonable expectation of success. See In re Merck & Co., Inc., 800 F.2d 1091, 231 USPQ 375 (Fed. Cir. 1986) (see MPEP § 2143.02). Therefore, it would have been obvious to a person of ordinary skill in the art to incorporate the teachings of Meade et al. with Holen et al. in order to provide for an effective detection method.

6. Claim 40 is rejected under 35 U.S.C. 103(a) as being unpatentable over Holen et al., Kearney et al. and Meade et al., as applied to claims 35, 36, 38 and 39 above, and further in view of Grinstaff et al. (U.S. Pat. No. 6,288,221 B1). Holen et al. and Mead et al. do not specifically

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teach the use of metallocenes. Grinstaff et al. do teach the use of a metallocene in methods for detecting nucleic acid sequences (see col. 9, lines 27 - 46; col. 42, lines 27 - 37). Hence, as evidenced by Grinstaff et al., a person of ordinary skill in the art would have recognized the use of metallocenes as a component of a detectable marker in a methodology for detecting target nucleic acid sequences. The Courts have held that the selection of a known material, based upon its suitability for the intended use, is within the ambit of one of ordinary skill in the art. See In re Leshin, 125 USPQ 416 (CCPA 1960) (see MPEP § 2144.07). Furthermore, since both Holen et al. and Grinstaff et al. rely on optical methods for detection, a person of ordinary skill in the art would accordingly have had a reasonable expectation of success of incorporating the use of a metallocene transition metal complex with the methods disclosed by Holen et al. (see Holen et al.: col. 19, line 65 – col. 20, line 38; Grinstaff et al.: col. 11, lines 26 – 65). The Courts have held that the prior art can be modified or combined to reject claims as prima facie obvious as long as there is a reasonable expectation of success. See In re Merck & Co., Inc., 800 F.2d 1091, 231 USPQ 375 (Fed. Cir. 1986) (see MPEP § 2143.02). Therefore, it would have been obvious to a person of ordinary skill in the art to incorporate the use of a metallocene transition metal complex, as taught by Grinstaff et al., with the methodology of Holen et al. and Meade et al., in order to facilitate the effective detection of target nucleic acid sequences.

Response to Arguments

Applicant's arguments with respect to claims 24 – 42 have been considered but are moot in view of the new ground(s) of rejection.

The examiner acknowledges the statement of common ownership, filed 7/2/2004, therefore obviating the rejection of claims 24 – 42 under 35 U.S.C §103(a) as being unpatentable

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over Bamdad et al. (U.S. Pat. No. 6,541,617 B1). The final rejection, mailed 3/29/2004, is withdrawn. Prosecution is reopened.

Conclusion

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Brian J. Sines, Ph.D. whose telephone number is (571) 272-1263. The examiner can normally be reached on Monday - Friday (11:30 AM - 8 PM EST).

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Jill A. Warden can be reached on (571) 272-1267. The fax phone number for the organization where this application or proceeding is assigned is 703-872-9306.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

Maureen M. Wallenhorst

MAUREEN M. WALLENHORST

PRIMARY EXAMINER

GROUP 1900